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Potentially inappropriate proton-pump inhibitor prescription in the general population: a claims-based retrospective time trend analysis

Muheim, Leander ; Signorell, Andri ; Markun, Stefan ; Chmiel, Corinne ; Neuner-Jehle, Stefan ; Blozik, Eva ; Ursprung, Pascale ; Rosemann, Thomas ; Senn, Oliver

Abstract: Background: Proton-pump inhibitors (PPI) are among the most prescribed drugs worldwide, and a large body of evidence raises concerns about their inappropriate use. Previous estimates of inappropriate use varied due to different definitions and study populations. Aims: We aimed to measure the population-based incidence and time trends of PPI and potentially inappropriate PPI prescriptions (PIPPI) with a novel method, continuously assessing excessive cumulative doses based on clinical practice guidelines. We also assessed association of patient characteristics with PPI prescriptions and PIPPI. Methods: This was an observational study based on a large insurance claims database of persons aged >18 years with continuous claims records of 12 months. The observation period was January 2012 to December 2017. We assessed the incidence and time trends of PPI prescriptions and PIPPI based on doses prescribed, defining 11.5 g of pantoprazole dose equivalents during any consecutive 365 days (average daily dose >31 mg) as inappropriate. Results: Among 1,726,491 eligible persons, the annual incidence of PPI prescriptions increased from 19.7% (2012) to 23.0% (2017), ($p = <0.001$), and the incidence of PIPPI increased from 4.8% (2013) to 6.4% (2017), ($p = <0.001$). Age, male gender, drugs with bleeding risk and multimorbidity were independent determinants of PIPPI ($p = <0.001$ for all). Conclusions: This study provides evidence that one of the most prescribed drug groups is commonly prescribed inappropriately in the general population and that this trend is increasing. Multimorbidity and drugs with bleeding risks were strong determinants of PIPPI. Addressing PPI prescriptions exceeding guideline recommendations could reduce polypharmacy and improve patient safety.

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Potentially inappropriate proton-pump inhibitor prescription in the general population: a claims-based retrospective time trend analysis

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Abstract

Background: Proton-pump inhibitors (PPI) are among the most prescribed drugs worldwide, and a large body of evidence raises concerns about their inappropriate use. Previous estimates of inappropriate use varied due to different definitions and study populations.

Aims: We aimed to measure the population-based incidence and time trends of PPI and potentially inappropriate PPI prescriptions (PIPP) with a novel method, continuously assessing excessive cumulative doses based on clinical practice guidelines. We also assessed association of patient characteristics with PPI prescriptions and PIPPI.

Methods: This was an observational study based on a large insurance claims database of persons aged >18 years with continuous claims records of ≥12 months. The observation period was January 2012 to December 2017. We assessed the incidence and time trends of PPI prescriptions and PIPPI based on doses prescribed, defining ≥11.5 g of pantoprazole dose equivalents during any consecutive 365 days (average daily dose >31 mg) as inappropriate.

Results: Among 1,726,491 eligible persons, the annual incidence of PPI prescriptions increased from 19.7% (2012) to 23.0% (2017), ($p < 0.001$), and the incidence of PIPPI increased from 4.8% (2013) to 6.4% (2017), ($p < 0.001$). Age, male gender, drugs with bleeding risk and multimorbidity were independent determinants of PIPPI ($p < 0.001$ for all).

Conclusions: This study provides evidence that one of the most prescribed drug groups is commonly prescribed inappropriately in the general population and that this trend is increasing. Multimorbidity and drugs with bleeding risks were strong determinants of PIPPI. Addressing PPI prescriptions exceeding guideline recommendations could reduce polypharmacy and improve patient safety.

Keywords: inappropriate, overuse, PPI, proton-pump inhibitors, polypharmacy

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Introduction

Proton-pump inhibitors (PPI) are highly effective antacid drugs¹ and are among the most commonly prescribed drugs worldwide. Pantoprazole, one of the most popular examples, is the single most prescribed generic drug and generates the highest overall generic drug cost in Switzerland.² As the use of PPI has been steadily increasing, concerns about their inappropriate use have been

growing. Both potential side effects^{3–5} and potentially unnecessary expenditures^{6–8} are of increasing interest. Many initiatives and professional societies advise against the prolonged use of PPI.^{9,10}

From a patient- and health-economics perspective, excessive average long-term doses seem most relevant, as higher cumulative doses of PPI might

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produce more side effects³ and cause more unnecessary costs.

Yet inappropriate use of PPI remains widespread and is still increasing, as multiple studies in various settings show.^{11–14} Many previous estimates of potentially inappropriate PPI prescriptions (PIPPI) are highly variable, but do not define inappropriateness based on continuous and cumulative doses prescribed on the patient level.^{11,13} Accordingly, considerable uncertainty remains about the valid extent of PIPPI within the population.

Aims

We aimed to measure the population-based incidence and time trends of PPI prescriptions and PIPPI by means of a novel method, which continuously assesses excessive cumulative doses based on clinical practice guidelines. We also assessed association of patient characteristics with PIPPI in comparison to PPI prescriptions.

Methods

Study design, data source and population

In this retrospective analysis, we used claims data of a major Swiss health insurance company ('Helsana group') covering approximately 14% of the Swiss population.¹⁵ All persons registered in Switzerland are obliged by law to contract basic health insurance. Individuals who cannot afford to pay the monthly premiums receive subsidies or full insurance coverage by their social services. Equal access to the full range of medical treatments including drugs is thereby granted to everyone, regardless of insurance status. Lower monthly premiums are available when choosing different deductible schemes. Deductibles range from 300 to 2500 Swiss Francs (CHF) yearly. In addition, optional alternative health plans (managed care, telemedicine gatekeeping and others) enable lower monthly premiums.

Our sample includes persons aged 18 or older whose records were available for at least 12 months from 1 January 2012 through 31 December 2017. Prescriptions of implausibly high cumulative doses were excluded.

Objectives

We defined our main objectives as follows:

1. **Incidence of PPI** prescriptions and time trends thereof (period 2012–2017)
2. **Incidence of PIPPI** and time trends thereof (period 2013–2017, PIPPI is defined as a result of the preceding year)
3. **Association of patient characteristics with PIPPI** (demographic characteristics, drugs with bleeding risk and number of chronic conditions) in comparison to PPI prescriptions

Assessment of proton-pump inhibitor prescribing

The insurance claims cover all pharmacy and physician invoices of prescription drugs. PPI were identified by Anatomical Therapeutic Chemical (ATC) Classification codes A02BC. Incidence of PPI prescriptions was defined by any such prescription during any calendar year. All health care invoices submitted to Helsana for reimbursement during the study period were considered. Prescribed PPI doses of the various PPI products were converted into *pantoprazole dose equivalents*, thus referencing the most commonly prescribed PPI product in Switzerland. In an attempt to estimate PPI prescribing in the context of clinical practice in Switzerland, doses of all the PPI agents on the market were labelled as either maintenance or therapeutic dose and then translated into pantoprazole dose equivalents of 20 mg and 40 mg, respectively. Detailed definitions of the various pantoprazole dose equivalents are listed under Supplemental Material (1).

Definition and assessment of potentially inappropriate proton-pump inhibitor prescribing

Based on a literature review covering 34 clinical practice guidelines in 2017, we extracted PPI dosage and treatment regimens on 57 different PPI indications.¹⁶ Thereof we derived a 365-day cumulative pantoprazole dose equivalent of 11.5 g as a cut-off for inappropriateness. This cut-off reflects an evidence-based maximal-dose scenario, based on three hypothetical sequential phases: (1) an 8-week double-dose therapeutic treatment phase (2×40 mg of pantoprazole dose equivalents, i.e. for severe erosive esophagitis),

(2) a 4-week tapering to a therapeutic dose (1×40 mg), (3) a maintenance dose (1×20 mg) for the remainder of the year. The resulting cumulative yearly dose of 11.2 g of pantoprazole equivalents was rounded up to 11.5 g. PIPPI was then determined by screening every observed person day for a preceding 365-day cumulative dose exceeding 11.5 g. To clarify we give a practical example: if 100 tablets pantoprazole 40 mg were dispensed 1 December 2016, this corresponded to 4000 mg added to the balance sheet of one patient. If the same patient repeated this process twice (i.e. adding another 8000 mg and therefore 12,000 mg in total) before 30 November 2017, then this patient would be regarded as a dispenser of PIPPI for 2017 (on the day of the third dispensing, i.e. when 11.5 g are exceeded).

Patient characteristics

Patient characteristics included gender, age and the number of chronic conditions according to ATC codes and pharmaceutical cost groups (PCG).¹⁷ Drugs which increase gastrointestinal bleeding risk and are therefore commonly associated with PPI co-prescriptions¹⁸ were assessed by translating the prescription of all such drugs into defined daily doses per patient per day, using corresponding ATC codes. These drugs were grouped as follows: Anticoagulant therapy including vitamin K antagonists (ATC B01AA) and direct oral anticoagulants (ATC B01AF, B01AE), antiplatelet therapy including aspirin (ATC B01AC06) and other platelet inhibitors (ATC B01AC), systemic corticosteroids (ATC H02AB) and non-steroidal anti-inflammatory drugs (NSAIDs), including coxibs (ATC M01A, M01AH). The number of chronic conditions using PCGs was likewise applied on the basis of drug prescriptions, according to all concurrent ATC codes billed (including ATC code A02 for acid-related disorders) per calendar year per patient, and categorized as follows: 0 (reference group), 1–3 and ≥ 4 .

Statistics: analysis

We calculated the incidence of PPI prescriptions, being the first PPI prescription during a calendar year and the incidence of PIPPI, being patients crossing at least once the cumulative prescribed PPI cut-off dose of 11.5 g during any preceding 365 days, counted per calendar year. Both measures used the total patients enrolled within the

calendar year as the denominator. In addition, the incidence of PIPPI among the incidental PPI prescription group was calculated. Percentages and mean values were used to describe categorical and nominal data, respectively. Descriptive statistics were used to provide characteristics of all patients with PPI prescriptions. These data were grouped into PPI prescriptions (i.e. non-PIPPI) and PIPPI groups. Chi-Squared tests for trend in proportions were used to analyse changes in proportions of PPI prescriptions and PIPPI incidence. For bivariate comparisons between the PPI prescriptions and PIPPI groups, Kruskal–Wallis, Fisher exact and Chi-Square tests were used accordingly. A two-sided *p*-value of 0.05 was considered statistically significant. To explore patient characteristics associated with PIPPI (dependent variable), we performed a multivariable logistic regression. Independent factors included in the regression model were age, gender, prescription of drugs increasing bleeding risk (yes/no) and number of chronic conditions. The goodness-of-fit of the logistic regression model was assessed by calculating the area under the curve (AUC).

To investigate a potential dose–response relationship between drugs increasing bleeding risk and PIPPI, we performed an additional analysis in which we included the absolute prescribed defined daily doses (DDDs)¹⁹ of any of the specific drug categories (dependent variables) into a regression model. For this purpose, we defined the corresponding time period in the group of PPI prescriptions as the 365 days' period following the first prescription of a PPI (as opposed to the retrospective 365 days in the PIPPI group). We present results of regression analysis as odds ratios with 95% confidence intervals (95% CI). Statistical analyses were conducted with “R” version 3.6.1 (2019-07-05).

Ethics

According to the Swiss national ethical and legal regulations, ethical approval from the local ethics committee was not needed for the study. As data were completely anonymized, no patient consent was necessary.

Results

The total amount of different persons accessible in our dataset throughout all years was 1,726,491.

Table 1. Annual incidence of PPI and PIPPI prescribing.

	2012	2013	2014	2015	2016	2017
Persons accessible (n)	1,197,992	1,188,017	1,181,640	1,195,286	1,169,965	1,105,847
Persons included (n)	939,295	941,897	933,377	933,450	917,590	870,073
PPI prescription incidence (n) (%)[*]	185,093 (19.7%)	199,477 (21.2%)	207,908 (22.3%)	214,448 (23.0%)	215,219 (23.5%)	200,565 (23.1%)
PIPPI incidence (n) (%)[*]	25,639 (2.7%)[†]	45,055 (4.8%)	50,418 (5.4%)	54,575 (5.8%)	56,504 (6.2%)	55,685 (6.4%)
Incidence of PIPPI among PPI prescriptions (%)	13.9%	22.6%	24.3%	25.5%	26.3%	27.8%

PIPPI, potentially inappropriate proton-pump inhibitors; PPI, Proton-pump inhibitors.
^{*} $p < 0.001$ for trend 2012–2017.
[†]Limited comparability of inception year inherent to definition of PIPPI.

After applying inclusion criteria, this number amounted to 1,337,722. The number of included persons is reported on a yearly basis in Table 1. The mean age during the observation period was 51.2 years, 52.4% of persons were female.

Incidence of PPI prescriptions and PIPPI, relative proportions in the study population and the group with PPI prescriptions are reported in Table 1. The incidence of PPI prescriptions was 23.0% in 2017, which reflects a steady rise since 2012 ($p < 0.001$ for trend). The overall incidence of PIPPI was 6.4% in 2017, corresponding to 27.8% of incident PPI prescriptions. Again, the positive trend for proportions since 2012 was statistically significant ($p < 0.001$ for trend).

All incident PPI prescriptions and PIPPI cases and corresponding patient characteristics are listed in Table 2. Patients with PIPPI were significantly older and more often female (unadjusted analysis).

The majority of patients with PPI prescriptions or PIPPI were also prescribed drugs with increased bleeding risk at least once during the designated 365-day periods, 63.3% and 86.0%, respectively. The highest incidence for a specific category of drugs with increased bleeding risk was observed for NSAIDs, with 50.9% and 62.0% of PPI prescriptions and PIPPI patients exposed.

Overall, the majority of patients in both prescription groups had at least one chronic condition: 73.2% of patients in the PPI prescriptions as compared with 98% in the PIPPI group. In the PPI prescriptions group, 1–3 chronic conditions were most prevalent. In the PIPPI group, the proportion of patients suffering from more than three chronic conditions reached 64%.

Results of the multivariable logistic regression of PIPPI and patient characteristics are presented in Table 3 and indicate that the model explains most of the variation of PIPPI (AUC 81.3%). In contrast to the bivariate comparison, female gender was independently associated with lower odds of PIPPI. All categories of concurrent drugs with bleeding risk and the number of chronic conditions remained independently associated with higher odds of PIPPI.

The additional analysis assessing the association of PIPPI with DDDs of all drugs with bleeding risk showed a significant dose–response relationship (Supplemental Material).

Discussion

In this large population-based study, one out of four adults was prescribed a PPI in 2017. Among these, one out of four was prescribed PIPPI based on cumulative doses not supported by clinical

Table 2. Patient characteristics and comparison between PPI and PIPPI prescribing groups.

	Characteristics			<i>p</i> -value for comparison
	Total	PPI	PIPPI	
<i>n</i>	610,437 (100%)	367,159 (60.1%)	243,278 (39.9%)	
Age (SD)	58.5 (19.6)	52.2 (19.8)	68.1 (14.9)	<0.001
Female gender (<i>n</i>) (%)	351,962 (57.7%)	209,852 (57.2%)	142,110 (58.4%)	<0.001
Drugs with bleeding risk (<i>n</i>) (%)				
Anticoagulants	68,557 (11.2%)	23,839 (6.5%)	44,718 (18.4%)	<0.001
Antiplatelets	138,998 (22.8%)	46,971 (12.8%)	92,027 (37.8%)	<0.001
Corticosteroids	151,595 (24.8%)	63,498 (17.3%)	88,097 (36.2%)	<0.001
NSAIDs	337,758 (55.3%)	186,908 (50.9%)	150,850 (62.0%)	<0.001
Any (of the above)	441,746 (72.4%)	232,573 (63.3%)	209,173 (86.0%)	<0.001
Number of chronic conditions (<i>n</i>) (%)				
0	100,474 (17.0%)	95,826 (26.8%)	4648 (2.0%)	<0.001
1–3	271,498 (45.9%)	191,706 (53.6%)	79,792 (34.1%)	<0.001
≥4	219,606 (37.1%)	69,810 (19.5%)	149,796 (64.0%)	<0.001
NSAIDs, non-steroidal anti-inflammatory drugs; PIPPI, potentially inappropriate proton-pump inhibitors; PPI, proton-pump inhibitors; SD, standard deviation.				

practice guidelines. We found increasing annual incidence trends for both PPI prescriptions and PIPPI during the observed 6-year period. Multimorbidity and drugs with bleeding risk were strong independent determinants of PIPPI.

In our study, the proportion of patients with at least one PPI prescription per year was 19.7% in 2012 and 23% in 2017. Corresponding rates from other health care systems varied between 7.4% and 29.8%.^{13,14,20–22} The definition of inappropriateness and differences between populations (i.e. inclusion criteria) and health care systems, including varying degrees of over-the-counter (OTC) dispensing, might explain some of this variance.

The annual incidence of PIPPI in our study rose from 4.8% in 2013 to 6.4% of all adults in 2017, corresponding to 27.8% of the population with an incidental PPI prescription. To our knowledge, there is no previous literature describing PIPPI

with continuous, cumulative dose regimens on the patient level and based on clinical practice guidelines. Multiple studies have assessed the extent of PIPPI in various settings and with different definitions of long-term use.¹² Previous estimates have been highly variable and ranged from proportions of overuse from 11 to 84%.²³ PIPPI was often ascertained either based on prescriptions exceeding a certain amount of weeks,^{24,25} within specific age groups,²⁵ specific situations such as drugs with bleeding risk,⁸ or as the absence of an appropriate clinical indication.^{14,26} A nationwide drug utilization study in France reported 32.4% of new PPI users in the year 2015 without other drugs or diagnoses to support an indication for PPI therapy,¹⁴ yet only 4.1% of new PPI users received a continuous PPI therapy lasting longer than 6 months. Another study assessed DDDs per patient but reported averages per calendar year, thereby leaving uncertainty about the actual degree of PIPPI at the patient level.¹¹ Nevertheless, some previous studies using different definitions of PIPPI,

Table 3. Adjusted association between PIPPI prescribing and patient characteristics.

Characteristics	OR	95% CI	p-value
Female gender	0.828	0.818–0.839	<0.001
Age (years)	1.027	1.027–1.027	<0.001
Drugs with bleeding risk			
Anticoagulants	1.376	1.350–1.403	<0.001
Antiplatelets	1.466	1.443–1.488	<0.001
Corticosteroids	1.557	1.536–1.580	<0.001
NSAIDs	1.154	1.139–1.169	<0.001
Number of chronic conditions (reference = 0)			
Chronic conditions (=1–3)	5.112	4.953–5.275	<0.001
Chronic conditions (≥ 4)	16.632	16.094–17.188	<0.001
AUC 81.3%			
AUC, area under the curve; 95% CI, 95% confidence intervals; NSAIDs, non-steroidal anti-inflammatory drugs; OR, odds ratio; PIPPI, potentially inappropriate proton-pump inhibitor.			

presented findings aligned with our results: a nationwide population-based study in Iceland found that 21% of incident high-dose PPI prescriptions were still using high doses after 1 year.¹³

Our data show a significant increase of PPI prescriptions and PIPPI over time. This phenomenon was described previously.^{13,20,27} Whether this is merely a consequence of an increased prevalence of polypharmacy in an aging society,²⁵ or pointing to challenges in the deprescribing process, with patients reporting increased symptoms and lower satisfaction after withdrawal of PPI,²⁸ remains unclear based on our data.

Higher age was associated with PIPPI compared with PPI prescriptions, whereas female gender was not. Yet evidence with regard to gender as determinant of PIPPI is conflicting.^{24,29}

Drugs with increased bleeding risk and especially NSAIDs were more common in the PIPPI group (62%), as compared with the PPI prescriptions group (50.9%). Similar proportions were reported elsewhere: NSAID co-use reached 48% in one study assessing long-term PPI prescriptions, and

a recent population-based estimate attributed NSAID use to 53.3% of patients newly prescribed a PPI.^{21,14} Our finding remained stable after regression analysis, showing that all drugs with bleeding risk were independently associated with PIPPI. The effect of NSAIDs decreased with short-term NSAID use, presumably due to a large number of healthy patients within the sample. While these data underline the substantial contribution of NSAIDs to PPI prescriptions, it remains unclear as to why exactly drugs with increased bleeding risk are associated to PIPPI (as opposed to PPI prescriptions). Interestingly, most guidelines commenting on prevention of NSAID-associated ulcers and bleedings (as compared with other guidelines recommending PPIs) leave room for interpretation by omitting recommendations on dose reduction in this specific context.^{30–33} As physicians tend to overestimate the need for preventive PPI prescriptions,³⁴ this might lead to prescriptions of higher doses more frequently. Also, it is known that previously asymptomatic patients can become symptomatic upon withdrawal of PPI drugs.³⁵ Patients with preventive PPI prescriptions might thus be at particular risk for the re-initiation of therapeutic doses, as

such prescriptions are often stopped abruptly in clinical practice, together with drugs increasing bleeding risk.

Furthermore, we found chronic conditions in 98% of patients with PIPPI and in 73.1% of patients with PPI prescribing. In fact, an increasing number of chronic conditions was the strongest predictor of PIPPI compared with PPI prescriptions in our logistic regression analysis. This finding has been described previously.^{21,25,36}

Strengths of this study are its large and representative sample including roughly 1.2 million persons, reflecting about 14% of the adult Swiss resident population.¹⁵ Furthermore, our method of defining PIPPI overcomes some of the variability of past estimates. It does not depend on the documentation and identification of (often absent) disease- or drug-related reasons justifying PPI prescriptions. It also does not rely on methods based on treatment intervals or days covered, which might not capture the long-term recurrence of PPI prescriptions or the actual dosage prescribed. In addition, by using a continuous, 'rolling' approach, our analysis is not restricted to calendar year boundaries and thereby can assess daily average doses over longer periods. The chosen cut-off value for PIPPI (11.5 g) is based on a clinical worst-case scenario of maximally necessary and sensible PPI intake, which only is surpassed in very seldom clinical situations.

This study has also limitations. We were not able to estimate OTC or hospital dispensing. However, available OTC formulations are low dose and small package size only, and patients are unlikely to use OTC PPI on a long-term basis, as costs are high and health insurance coverage is plenary. Furthermore, the effective ingestion of PPI of course remains unclear, as our data documents drug reimbursement only. Again, it is unlikely that a relevant amount of patients would regularly dispense PPI on a long-term basis without actually ingesting them. Lastly, our sample does not contain clinical data. We therefore cannot exclude potentially justified long-term, high-dose PPI treatments as occurs in very rare conditions. However, clinical recommendations for long-term prescriptions of therapeutic dose PPI are limited: Barrett-esophagus³⁷ with a prevalence of <2% in the general adult population³⁸ and Zollinger–Ellison syndrome, which is exceedingly rare.³⁹ It seems thus unlikely, that such disease-specific

therapeutic dose PPI prescriptions would alter the general finding of this study, which affects a substantial share of the overall population.

Many studies observed side effects of PPI prescriptions use without establishing causality,⁴⁰ but some trials did.^{41,42} In light of the widespread prescription of PPI, even the probability of rare or minor side effects should raise concern. Appropriate PPI prescriptions (and deprescribing) in the context of short-term preventive use (e.g. with NSAIDs) might be of special interest for further investigations. Co-occurrence is widespread and specific recommendations on dose reductions are lacking. Our data portray a period when dose lowering and deprescribing of PPI should have been commonplace. The *Choosing Wisely* campaign in the US adopted this target in 2012,¹⁰ which was widely received in Switzerland. Its Swiss offshoot *Smarter Medicine*⁴³ absorbed the PPI lowering recommendation later, in 2015.

However, a systematic review reported PPI to be among the drugs most resistant to deprescribing.⁴⁴ A call for more deprescribing⁴⁵ might therefore not be enough. Some authors, as a consequence, believe that prescribing or paying restrictions (self-pay for high dose) should be put in place by authorities.^{21,45} We suggest that a standardized operationalization of PIPPI, as presented in this study, offers opportunities to measure PIPPI use more validly, to compare its occurrence across health service providers, settings, and care models, and possibly to incentivize a more cautious use of PPI drugs, while increasing transparency and awareness.

Conclusion

This study provides evidence that one of the most prescribed drug groups is commonly prescribed inappropriately in the general population and that this trend is increasing. Multimorbidity and drugs with bleeding risks were strong determinants of PIPPI. Addressing PPI prescriptions exceeding guideline recommendations could reduce polypharmacy and improve patient safety.

Author contributions

All authors have been involved in the study concept, design and interpretation of results. Andri Signorell collected and analysed the data, Leander Muheim drafted the article and all of the authors revised it critically. All authors

approved the final version of the article, including the authorship list.

Conflict of interest statement

Helsana Group provided support in the form of salaries for authors AS and EB being employed by the Helsana Group. The funders had no role in the study design, data collection and analysis, decision to publish, or preparation of the manuscript. AS and EB declare no conflict of interest. Helsana Group shall have no liability to any third party in respect of the contents of this article. All the other authors as well have no conflicts of interests or financial disclosures to declare.

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Supplemental material

Supplemental material for this article is available online.

References

1. Katz PO, Gerson LB and Vela MF. Guidelines for the diagnosis and management of gastroesophageal reflux disease. *Am J Gastroenterol* 2013; 108: 308–328; quiz 329.
2. Schneider R, Reinau D, Schur N, *et al.* Helsana-Arzneimittel. Report, Universität Basel, Ausgabe, 2018, p.250.
3. Poly TN, Islam MM, Yang H-C, *et al.* Proton pump inhibitors and risk of hip fracture: a meta-analysis of observational studies. *Osteoporos Int* 2019; 30: 103–114.
4. Xie Y, Bowe B, Li T, *et al.* Risk of death among users of proton pump inhibitors: a longitudinal observational cohort study of United States veterans. *BMJ Open* 2017; 7: e015735.
5. Makunts T, Cohen IV, Awdishu L, *et al.* Analysis of postmarketing safety data for proton-pump inhibitors reveals increased propensity for renal injury, electrolyte abnormalities, and nephrolithiasis. *Sci Rep* 2019; 9: 2282.
6. Harrison SL, Kouladjian O'Donnell L, Milte R, *et al.* Costs of potentially inappropriate medication use in residential aged care facilities. *BMC Geriatr* 2018; 18: 9.
7. Ladd AM, Panagopoulos G, Cohen J, *et al.* Potential costs of inappropriate use of proton pump inhibitors. *Am J Med Sci* 2014; 347: 446–451.
8. Chau SH, Sluiter RL, Hugtenburg JG, *et al.* Cost-utility and budget impact analysis for stopping the inappropriate use of proton pump inhibitors after cessation of NSAID or low-dose acetylsalicylic acid treatment. *Drugs Aging* 2020; 37: 67–74.
9. Kahrilas PJ, Shaheen NJ and Vaezi MF. American gastroenterological association medical position statement on the management of gastroesophageal reflux disease. *Gastroenterology* 2008; 135: 1383–1391.e5.
10. Treating GERD. American Gastroenterological Association. Choosing Wisely, <http://www.choosingwisely.org/clinician-lists/american-gastroenterological-association-treating-gerd/> (accessed 12 March 2019).
11. Haastrup P, Paulsen MS, Zwisler JE, *et al.* Rapidly increasing prescribing of proton pump inhibitors in primary care despite interventions: a nationwide observational study. *Eur J Gen Pract* 2014; 20: 290–293.
12. Raghunath AS, O'Morain C and McLoughlin RC. Review article: the long-term use of proton-pump inhibitors. *Aliment Pharmacol Ther* 2005; 22(Suppl. 1): 55–63.
13. Hálfðánarson ÓÖ, Pottegård A, Björnsson ES, *et al.* Proton-pump inhibitors among adults: a nationwide drug-utilization study. *Therap Adv Gastroenterol* 2018; 11: 1756284818777943.
14. Lassalle M, Le Tri T, Bardou M, *et al.* Use of proton pump inhibitors in adults in France: a nationwide drug utilization study. *Eur J Clin Pharmacol* 2020; 76: 449–457.
15. Tabelle. Bilanz der ständigen Wohnbevölkerung, 1861–2018. Bundesamt für Statistik, <https://www.bfs.admin.ch/bfs/de/home/statistiken/bevoelkerung/stand-entwicklung/bevoelkerung.assetdetail.13707405.html> (2019, accessed 21 June 2020).
16. Ursprung P. *Protonenpumpenhemmer in klinischen guidelines*. Master Thesis, Medical Faculty University of Zurich, Switzerland, 2020.

17. Huber CA, Szucs TD, Rapold R, *et al.* Identifying patients with chronic conditions using pharmacy data in Switzerland: an updated mapping approach to the classification of medications. *BMC Public Health* 2013; 13: 1030.
18. Lin KJ, Hernández-Díaz S and García Rodríguez LA. Acid suppressants reduce risk of gastrointestinal bleeding in patients on antithrombotic or anti-inflammatory therapy. *Gastroenterology* 2011; 141: 71–79.
19. WHO Collaborating Centre for Drug Statistics Methodology. ATC/DDD index, https://www.whocc.no/atc_ddd_index/ (accessed 21 August 2018).
20. Othman F, Card TR and Crooks CJ. Proton pump inhibitor prescribing patterns in the UK: a primary care database study. *Pharmacoepidemiol Drug Saf* 2016; 25: 1079–1087.
21. Daniels B, Pearson S-A, Buckley NA, *et al.* Long-term use of proton-pump inhibitors: whole-of-population patterns in Australia 2013–2016. *Therap Adv Gastroenterol* 2020; 13: 1756284820913743.
22. Pottegård A, Broe A, Hallas J, *et al.* Use of proton-pump inhibitors among adults: a Danish nationwide drug utilization study. *Therap Adv Gastroenterol* 2016; 9: 671–678.
23. Naunton M, Peterson GM, Deeks LS, *et al.* We have had a gutful: the need for deprescribing proton pump inhibitors. *J Clin Pharm Ther* 2018; 43: 65–72.
24. Moriarty F, Bennett K, Cahir C, *et al.* Characterizing potentially inappropriate prescribing of proton pump inhibitors in older people in primary care in Ireland from 1997 to 2012. *J Am Geriatr Soc* 2016; 64: e291–e296.
25. Cahir C, Fahey T, Teeling M, *et al.* Potentially inappropriate prescribing and cost outcomes for older people: a national population study. *Br J Clin Pharmacol* 2010; 69: 543–552.
26. Levy-Neumand O, Carniaux F, Bonaz B, *et al.* Proton pump inhibitors in general medicine. Comparison of routine practices with marketing authorization indications. *Gastroenterol Clin Biol* 2007; 31: 78–83.
27. Haastrup PF, Paulsen MS, Christensen RD, *et al.* Medical and non-medical predictors of initiating long-term use of proton pump inhibitors: a nationwide cohort study of first-time users during a 10-year period. *Aliment Pharmacol Ther* 2016; 44: 78–87.
28. Boghossian TA, Rashid FJ, Thompson W, *et al.* Deprescribing versus continuation of chronic proton pump inhibitor use in adults. *Cochrane Database Syst Rev* 2017; 3: CD011969.
29. Wallerstedt SM, Fastbom J, Linke J, *et al.* Long-term use of proton pump inhibitors and prevalence of disease- and drug-related reasons for gastroprotection—a cross-sectional population-based study. *Pharmacoepidemiol Drug Saf* 2017; 26: 9–16.
30. Laine L and Jensen DM. Management of patients with ulcer bleeding. *Am J Gastroenterol* 2012; 107: 345–360.
31. Malfertheiner P, Megraud F, O’Morain CA, *et al.* Management of *Helicobacter pylori* infection—the Maastricht IV/ Florence Consensus Report. *Gut* 2012; 61: 646–664.
32. Satoh K, Yoshino J, Akamatsu T, *et al.* Evidence-based clinical practice guidelines for peptic ulcer disease 2015. *J Gastroenterol* 2016; 51: 177–194.
33. Fischbach W, Malfertheiner P, Jansen PL, *et al.* S2k-leitlinie *helicobacter pylori* und gastroduodenale Ulkuserkrankheit. *Z Gastroenterol* 2016; 54: 327–363.
34. Murthy SK, Kauldher S and Targownik LE. Physicians’ approaches to the use of gastroprotective strategies in low-risk non-steroidal anti-inflammatory drug users. *Aliment Pharmacol Ther* 2006; 23: 1365–1372.
35. Reimer C, Søndergaard B, Hilsted L, *et al.* Proton-pump inhibitor therapy induces acid-related symptoms in healthy volunteers after withdrawal of therapy. *Gastroenterology* 2009; 137: 80–87, 87.e1.
36. Voukelatou P, Vrettos I, Emmanouilidou G, *et al.* Predictors of inappropriate proton pump inhibitors use in elderly patients. *Curr Gerontol Geriatr Res* 2019; 2019: 7591045.
37. Jankowski JAZ, de Caestecker J, Love SB, *et al.* Esomeprazole and aspirin in Barrett’s oesophagus (AsPECT): a randomised factorial trial. *Lancet* 2018; 392: 400–408.
38. Ronkainen J, Aro P, Storskrubb T, *et al.* Prevalence of Barrett’s esophagus in the general population: an endoscopic study. *Gastroenterology* 2005; 129: 1825–1831.
39. Öberg K and Eriksson B. Endocrine tumours of the pancreas. *Best Pract Res Clin Gastroenterol* 2005; 19: 753–781.
40. Yadlapati R and Kahrilas PJ. When is proton pump inhibitor use appropriate? *BMC Med* 2017; 15: 36.

41. Moayyedi P, Eikelboom JW, Bosch J, *et al.* Safety of proton pump inhibitors based on a large, multi-year, randomized trial of patients receiving rivaroxaban or aspirin. *Gastroenterology* 2019; 157: 682–691.e2.
42. Writing Committee for the American Lung Association Asthma Clinical Research Centers, Holbrook JT, Wise RA, *et al.* Lansoprazole for children with poorly controlled asthma: a randomized controlled trial. *JAMA* 2012; 307: 373–381.
43. smarter medicine - Choosing Wisely Switzerland. Top-5-Listen – Ambulante Allgemeine Innere Medizin, <https://www.smartermedicine.ch/> (accessed 20 January 2021).
44. Dills H, Shah K, Messinger-Rapport B, *et al.* Deprescribing medications for chronic diseases management in primary care settings: a systematic review of randomized controlled trials. *J Am Med Dir Assoc* 2018; 19: 923–935.e2.
45. Henderson J, Bouck Z, Holleman R, *et al.* Comparison of payment changes and choosing wisely recommendations for use of low-value laboratory tests in the United States and Canada. *JAMA Intern Med* 2020; 180: 524–531.

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